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(54) **STEROIDAL SAPONIN PHARMACEUTICAL COMPOSITION, ITS PREPARATION METHOD AND USE**

(57) The present invention provides a pharmaceutical composition containing steroidal saponins. The pharmaceutical composition comprises 5~25 parts by weight of furostanal saponins represented by general formula A or general formula B and 1~10 parts by weight of spirostanol saponin represented by general formula C. The present invention further provides a method for preparing the pharmaceutical composition. After determination of the three steroidal saponin components in the pharma-

ceutical composition of the present invention, it is proved that it has high stability and liability in its therapeutic effect. Meanwhile, the method to identify the three steroidal saponins is provided with reliable controllability. Besides, with its daily dosage of 300~600 mg, which is less dosage, the drug of this invention provides a new choice in clinics.

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Description**Technical Field of the Invention**

[0001] The present invention relates to a pharmaceutical composition containing steroidal saponins, particularly, relates to a pharmaceutical composition containing steroidal saponins mainly extracted from the raw materials of *Dioscorea panthaica* Prain et Burkill and *Dioscorea nipponica* Makino.

Background of the Invention

[0002] Steroidal saponins are a sort of important bioactive substance in plants. The study of steroidal saponin has been occupying an important position in chemistry of natural products. The aglycon of steroidal saponin is spirostanol or furostanol containing 27 carbon atoms, and exists mostly in such monocotyledonous plants as Liliaceae, Amaryllidaceae and Dioscoreaceae.

[0003] The herb *Dioscorea panthaica* Prain et Burkill (a species in *Dioscorea*, *Dioscoreaceae*) is the rootstocks of herbal plant *Dioscorea panthaica* Prain et Burkill and has many other names such as curcuma, curcuma longae, tendrilleaf solomonseal rhizome and Turmeric Root Tuber. It is distributed in Southwest China and Hubei and Hunan provinces of the country and has the actions of regulating *qi* (*vital energy*) to alleviate pain and detumescence by detoxification, so it is often prescribed for peratodynia, vomiting, diarrhea, abdominal pains, injuries, fractures, pyocutaneous diseases and venomous snake bites [Chinese Materia Medica, Vol 8 p8.247, compiled by Chinese Materia Medica Editorial Committee, State Traditional Chinese Medical Administration Bureau, and published by Shanghai Science and Technology Press in 1999]. The herb *Dioscorea nipponica* Makino (a species in *Dioscorea*, *Dioscoreaceae*) is the rootstocks of the herbal plant *Dioscorea nipponica* Makino and also owns many other names such as curcuma, monkshood vine root-bark and so on. It is distributed in North, Northeast and Northwest of China and also in Henan, Hubei, Shandong, Jiangsu, Anhui, Zhejiang, Jiangxi and Sichuan provinces of the country. It has the actions of expelling rheumatism and removing wetness, promoting blood circulation to remove meridian obstruction and stopping coughing, so it is often prescribed for rheumatic arthralgia, limb numbness, thoracic obstruction, cardiodynia, chronic tracheitis, injuries, fractures, malaria, anthracias and swelling [Chinese Materia Medica, Vol 8 p238, compiled by Chinese Materia Medica Editorial Committee, State Traditional Chinese Medical Administration Bureau, and published by Shanghai Science and Technology Press in 1999]. *Dioscorea nipponica* Makino was also listed on P435-436, Part I, Edition 1977 of Pharmacopoeia of the People's Republic of China.

[0004] Presently, researches on *Dioscorea panthaica* Prain et Burkill and *Dioscorea nipponica* Makino are rare. It has been reported by Bogang Li *et al.* that two kinds of water-insoluble steroidal saponins, dioscin and gracillin, were separated and identified from *Dioscorea panthaica* Prain et Burkill. [Bogang Li *et al.* ACTA BOTANICA SINICA 1986, 28(4): 409-411]; Three kinds of steroidal saponins were separated by Mei Dong *et al.* from the *Dioscorea panthaica* Prain et Burkill, the structures of which have been identified as pseudoprosteroidal saponins (Dioscin 1) and DI-9 (Dioscin 6) [Mei Dong *et al.* ACTA MEDICA SINICA 2001, 36(1): 42-45]. It is also reported that two kinds of water-insoluble steroidal saponins, dioscin and gracillin both being the components of perhexiline, were obtained and identified by Yiwei Fang *et al.* from *Dioscorea nipponica* Makino [Yiwei Fang *et al.* ACTA MEDICA SINICA 1982, 17(5): 388-391]; Two kinds of steroidal saponins were also extracted and obtained from *Dioscorea nipponica* Makino, in which one is water-insoluble and another is water-soluble, the structures of which are determined as gracillins [Shuhu Du *et al.* ACTA MEDICA SINICA 2002, 37(4): 267-270]. Another research discloses the structural formulas of eight kinds of steroidal saponins extracted from the total dioscins [Bogang Li; Zhengzhi Zhou. Traditional Chinese Drug Research & Clinical Pharmacology 1994, 13(2): 75-76]. Said eight kinds of steroidal saponins are the active components for treating coronary artery disease, including dioscins, protodioscins, protogracillins and gracillins. Chinese patent application CN020128119.X discloses a new use of six kinds of spirostanol steroidal saponins for treating Cardiovascular Diseases. Chinese patent application CN02112159.1 discloses a new use of a composition containing diosgenins for treating Myocardial Ischemia, Angina and Myocardial Infarction. Until now, only two kinds of components, pseudoprotodioscin (I) and dioscin (II), have been separated and identified from the herbs *Dioscorea panthaica* Prain et Burkill and *Dioscorea nipponica* Makino.

[0005] Although the above-mentioned references disclose that *Dioscorea panthaica* Prain et Burkill and *Dioscorea nipponica* Makino contain several kinds of steroidal saponins that are effective for treatment of cardiovascular diseases, the researches only relate to chemical studies on effective components of above two plants, and some of the components have little value in pharmaceuticals due to their low contents in the plants. As everyone knows, it would be very difficult and high cost to obtain effective and pure chemicals from plants. Meanwhile, in order to avoid unexpected side effects and to reduce costs and use conveniently, it is a general practice not to use pure effective compounds of herbal plants but use the extracts of the herbal plants as raw materials of pharmaceuticals. One disadvantage for such practice is that differences of seasons for harvesting the plants and growth areas thereof would result in changes in contents of effective

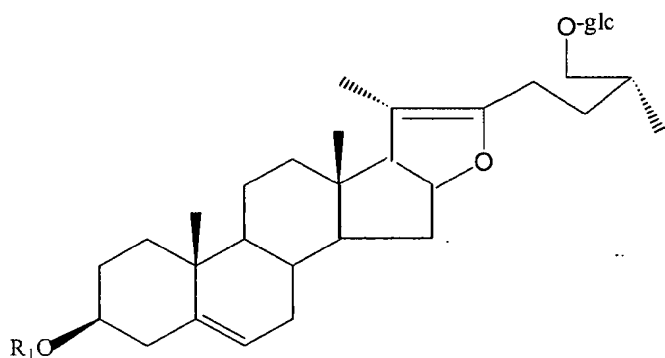
components. However, unclearness of main effective components and the contents thereof would make great difficulties in quality control of pharmaceuticals. Therefore, in order to maintain stability of quality of herbal medicine, it is important to get a clear idea on the basic components and contents of herbal plants. Due to technical limitation and complexity in species in steroidal saponins, there is no definitive report on main effective components of steroidal saponins, and the contents and purity thereof so far.

[0006] The traditional method for preparation of total steroidal saponins is as follows: after heat extraction with methanol or ethanol, condensing the alcoholic solution obtained; after removing methanol or ethanol, degreasing the solution with chloroform, extracting with normal butyl alcohol containing water. Total steroidal saponins are achieved after condensing the solution. The process described above has the following shortcomings: 1) low yield and high cost; 2) long production period; 3) in need of great amount of organic solvents, which might be risky of flaming or poisoning in production and bring about serious environmental pollution; 4) great energy consumption resulted from condensation and desiccation of the normal butyl alcohol with a high boiling point; 5) poorness in color and gloss of products and improperness for industrialized production. Total steroidal saponins prepared with traditional techniques are instable in contents, which would cause unstableness in effects of the drugs made thereof.

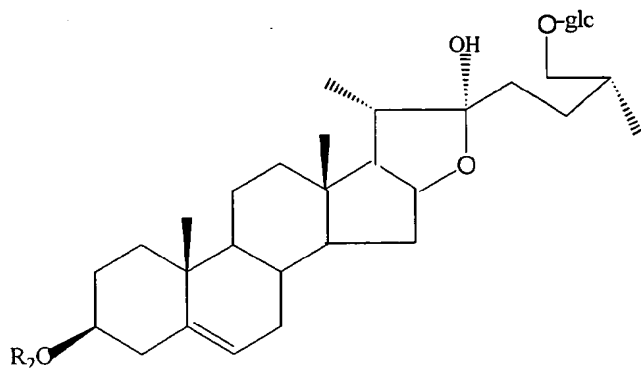
Summary of the Invention

[0007] In order to overcome the above shortcomings, the present invention provides a pharmaceutical composition containing steroidal saponins. The present invention also provides a method for preparing the same and a use thereof.

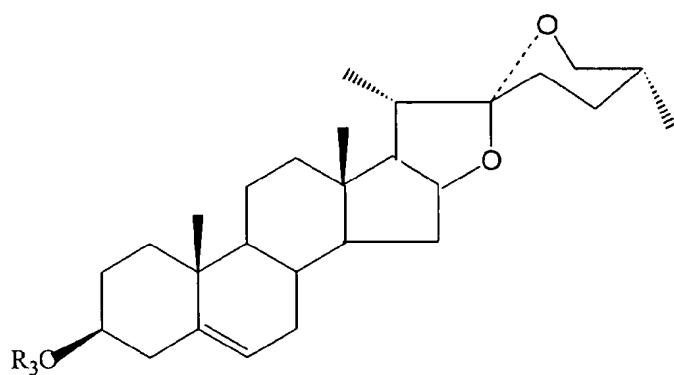
[0008] The present invention provides a pharmaceutical composition containing steroidal saponins, it comprises 5~25 parts by weight of furostanol saponin represented by general formula A and/or general formula B, and 1~10 parts by weight of spirostanol saponin represented by general formula C.



General Formula A

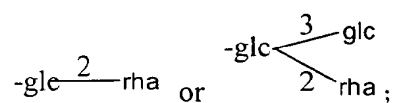
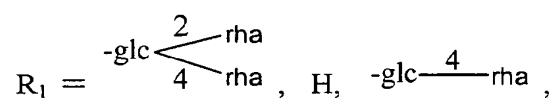


General Formula B

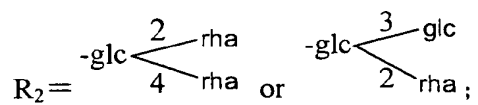


General Formula C

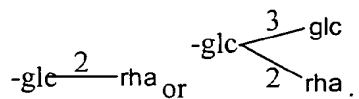
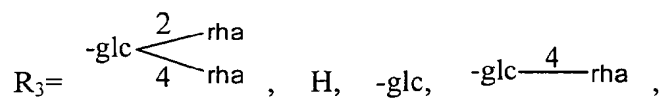
wherein, in general formula A,



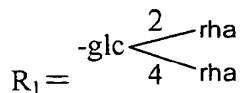
in general formula B:



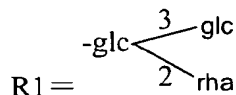
in general formula C:



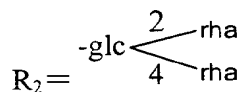
[0009] More preferably, it comprises following compounds: in general formula A:
when



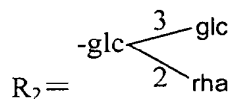
the compound represented by general formula A is pseudoprotodioscin (I);
when



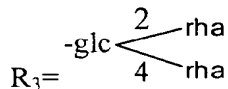
the compound represented by general formula A is pseudoprotogracillin (II); in general formula B,
when



the compound represented by general formula B is protodioscin (IV)
when



the compound represented by general formula B is protogracillin (V);
in general formula C, when



the compound represented by general formula C is dioscin (III),

[0010] In an embodiment, the composition of the present invention comprises 5~22 parts of pseudoprotodioscin (I) and/or protodioscin (IV), 1~5 parts of pseudoprotogracillin (II) and/or protogracillin (V) and 1~8 parts of dioscin (III).

[0011] Preferably, it comprises 5~20 parts of pseudoprotodioscin (I), 1~3 parts of pseudoprotogracillin (II) and 1~5 parts of dioscin (III).

[0012] More preferably, it comprises 12~18 parts of pseudoprotodioscin (I), 1 part of pseudoprotogracillin (II) and 1.2~2.5 parts of dioscin (III).

[0013] The steroidal saponins used in the composition are originated from the extracts of *Dioscorea panthaica* Prain et Burkill and *Dioscorea nipponica* Makino, both belonging to species of *Dioscorea*, *Dioscoreaceae*.

[0014] Wherein, in the said extract of total steroidal saponins, the content of total steroidal saponins is more than 80% (w/w), and no less than 65% (w/w) when calculated by dioscin.

[0015] Wherein, in the said extract, the total content of three kinds of steroidal saponins, i.e. pseudoprotodioscin (I), pseudoprotogracillin (II) and dioscin (III), is no less than 50% (w/w) of the content of total steroidal saponins.

[0016] The said extract has the HPLC fingerprint as shown in Fig 1, wherein the characteristic peaks of the fingerprint are as follows respectively: the retention time for pseudoprotodioscin (I) is 28.27 min; the retention time for pseudoprotogracillin (II) is 29.5 min and the retention time for dioscin (I) is 57.10 min.

[0017] Chromatographic condition: chromatographic column: Alltima C18 4.6×250mm, 5 m; Gradient elution; Deter-

mined with an evaporative light scattering detector; Drift tube temperature: 100°C; Gas flow rate: 2.0L / min

[0018] The said extract is prepared by the methods as follows:

a. Taking rootstocks of *Dioscorea panthaica* Prain et Burkill and *Dioscorea nipponica* Makino, or the rootstocks of freshly-harvested herbal plants *Dioscorea panthaica* Prain et Burkill and *Dioscorea nipponica* Makino as raw material. Extracting with mixed solution containing water, methanol, ethanol, n-butanol or mixed solution comprising one or more than one kinds of other low-fat alcohols after crushing and slicing the rootstocks, the amount of said mixed solution shall be 15~25 times of the raw material.

b. Cooling and filtering the extract prepared in step a, passing the filtrate through absorbent resin column, abandoning the effluent and washing with water till the effluent turns colorless, abandoning the rinsing water.

c. Eluting the absorbent resin column that is washed with water in step b with one or more than one kinds of solvent selected from a group consisting of ethanol, methanol, acetone, 50%~90% ethanol, 30~80% hydrous methanol and 60~95% hydrous acetone; collecting and condensing the effluent.

d. adding 60~95% alcohol to the concentrated solution obtained in step c for alcohol precipitation, filtering and collecting filtrate, and the product is obtained after condensing and drying the filtrate.

[0019] Wherein, in step b of the above method, it further includes a step of decompressing condensation before cooling and filtering when the extract contains methanol, ethanol, n-butanol or other low-fat alcohols.

[0020] The solvent used in step c can be one or more than one kinds of solvent selected from water, methanol, ethanol, n-butanol or other low-fat alcohols or mixture thereof. The extraction is conducted by soakage or ultrasonic resonance at room temperature, or by soakage or reflux under heating condition. The times for extraction might be once or several times.

[0021] The resins used in the step b can be one or more than one kinds of resins selected from a group consisting of HPD₁₀₀, HPD₃₀₀, LD₄₀, D₁₀₁, and mixture thereof. The solvent for elution can be one or more than one kinds of solvents selected from a group consisting of water, methanol, ethanol, acetone, hydrous methanol, hydrous ethanol and hydrous acetone. Elution can be either concentration elution or gradient elution.

[0022] The pharmaceutical composition of the present invention comprises steroidal saponins or the extracts thereof as an active ingredient and pharmaceutically acceptable adjuvants.

[0023] Wherein, the pharmaceutical composition of the present invention can be prepared to various forms, such as tablet, capsule, soft capsule, grain, oral liquor, dripping pill and injection.

[0024] The present invention also provides a method for preparing the pharmaceutical composition, including the following steps:

[0025] Taking by weighing furostanol saponins of general formula A and/or general formula B and spirostanol saponin of general formula C and mixing them at the ratio of 5~25 parts by weight of furostanol saponin and 1~5 parts by weight of spirostanol saponin. Adding pharmaceutically acceptable adjuvants, it could be then prepared to pharmaceutical composition.

[0026] Further, the pharmaceutical composition provided by the present invention can also be prepared by the following method:

a. Taking rootstocks of *Dioscorea panthaica* Prain et Burkill and *Dioscorea nipponica* Makino, or the rootstocks of freshly-collected herbal plants of the same as the raw material. After slicing or smashing the rootstocks, extracting with one or more than one kinds of solvent selected from a group consisting of water, methanol, ethanol, n-butanol and other low-fat alcohols, the amount of the solvent shall be 15~25 times of the raw material.

b. Cooling and filtering the extract prepared in step a, passing the filtrate through absorbent resin column, abandoning the effluent and washing with water till the effluent turns colorless, abandoning the rinsing water.

c. Eluting the absorbent resin column that was washed with water in step b with one or more than one kinds of solvent selected from a group consisting of ethanol, methanol, acetone, 50%~90% ethanol, 30~80% hydrous methanol and 60~95% hydrous acetone, collecting and condensing the effluent.

d. Adding 60~95% alcohol to the concentrated solution obtained in step c for alcohol precipitation, filtering and collecting the filtrate.

e. Condensing and drying the filtrate in step d, and adding pharmaceutically acceptable adjuvants to obtain the pharmaceutical composition in the forms of common usage.

[0027] Wherein, in step b, it further includes a step of decompressing condensation before cooling and filtering when the extract contains methanol, ethanol, n-butanol or other low-fat alcohols.

[0028] The present invention also provides a use of the pharmaceutical composition in preparing the pharmaceutical for treating and preventing cerebrocardiovascular diseases. Wherein, said pharmaceutical refers to that to treat coronary artery disease, angina, myocardial infarction, arrhythmia, hyperlipemia or ischemic cerebrovascular disease.

[0029] After determination of the three steroidal saponin components in the pharmaceutical composition of the present invention, it is proved that it has high stability and liability in its therapeutic effect. Meanwhile, the method to identify the three steroidal saponins is provided with reliable controllability. Besides, with its daily dosage of 300~600 mg, which is less dosage, the drug of this invention provides a new choice in clinics.

[0030] Obviously, according to the above-mentioned content of the present invention and by means of common knowledge and routine measures adopted in this field of technology, it is possible to make many amendments, replacements or alternations to the present invention under the prerequisite not to break away the basic ideas of the techniques mentioned above.

[0031] Following embodiments are used to further illustrate the present invention, but this by no means shall be understood as the limitation for the scope of the subjects of the present invention. Any technology realized on basis of the said content of the present invention belongs to the scope of the present invention.

Brief Description of Drawings

[0032]

FIG 1 is the HPLC spectrogram for protodioscin sample and the mixture of compounds 1~8, wherein peaks numbered 1~8 are correspondent to compounds 1~8;

FIG 2 is the HPLC spectrogram for the composition in example 1, wherein the peak No 1 is correspondent to compound 1, the peak No 6 is correspondent to compound 2, the peak No 7 is correspondent to compound 9, the peak No 8 is correspondent to compound 3 and the peak No 11 is correspondent to compound 5.

FIG 3 is the HPLC spectrogram for the composition in example 2, wherein the peak No 1 is correspondent to compound 1, the peak No 6 is correspondent to compound 2, the peak No 7 is correspondent to compound 9, the peak No 8 is correspondent to compound 3 and the peak No 10 is correspondent to compound 5.

FIG 4 is the HPLC spectrogram for the composition in example 3, wherein the peak No 1 is correspondent to compound 1, the peak No 6 is correspondent to compound 2, the peak No 7 is correspondent to compound 9, the peak No 8 is correspondent to compound 3 and the peak No 10 is correspondent to compound 5.

Detailed Description of the Invention

[0033] Example 1 Preparation of the pharmaceutical composition of the present invention:

100 kg dried rootstocks of *Dioscorea nipponica* Makino were taken and crushed. Reflux was made with 95% ethanol (1200L×3) for 3 hrs for the first time and then 2 hrs for the second and third times, respectively. After filtration, the extract was gathered and ethanol recovered. With addition of water to 2g / ml, it was refrigerated for 24 hrs before being centrifuged. The supernatant liquid was passed through HPD₃₀₀ resin column of adsorption, which was then washed with water till the effluent turned colorless before being eluted with 70% ethanol. The part eluted with 70% ethanol was collected and condensed. The concentrated solution was added with 75 % ethanol for precipitation before being filtrated. The filtrate was collected, condensed and dried. After being desiccated by vacuum dehydration, the product of total steroidal saponins of *Dioscorea nipponica* Makino was obtained. The yield was 2.32% and the content of total steroidal saponins was 85 % (w / w).

[0034] Example 2 Preparation of the pharmaceutical composition of the present invention:

100 kg dried rootstocks of herb *Dioscorea panthaica* Prain et Burkill were taken and sliced. They were boiled for 4 times with addition of 12 times of water. The time for first decoction is 3 hrs and then 2 hrs for the second, third and fourth times, respectively. After filtration, the extract was gathered and placed till it turned to room temperature. After being centrifuged, the supernatant liquid was passed through macroporous adsorptive resin column [HPD₁₀₀ / LD₁₀₀=7:3 (W / W)], which was then washed with water till the effluent turned colorless before being eluted with 10 % ethanol. Elution with 75 % ethanol were done finally and the part eluted with 75 % ethanol was collected and condensed. The concentrated solution was added with 80 % ethanol for precipitation before be filtrated. The filtrate was collected, condensed and dried till the filtrate is odorless after ethanol is recovered. After being desiccated again with the method of spray drying, the product of total steroidal saponins of *Dioscorea panthaica* Prain et Burkill was obtained. The yield was 1.5% and the content of total steroidal saponins was 90 % (w/w).

[0035] Example 3 Preparation of the pharmaceutical composition of the present invention:

400 kg dry rootstocks of fresh *Dioscorea nipponica* Makino were taken and sliced. They were boiled for 3 times with

addition of 8 times of water. The time for first decoction is 3 hrs and then 2 hrs for the second and third times, respectively. After filtration, the extract was gathered and placed till it cooled down to room temperature. After being centrifuged, the supernatant liquid was passed through macroporous adsorptive resin column [HPD₁₀₀ / LD₁₀₀=6:4(w/w)], which was then flushed and back flushed thoroughly with water before being eluted with 10% ethanol. Elution with 80 % ethanol were done finally and the part eluted with 80 % ethanol was collected and condensed. The concentrated solution was added with 90% ethanol for alcohol precipitation before being filtrated. The filtrate was collected, condensed, and dried till the filtrate is odorless after ethanol is recovered. After being desiccated again with the method of spray drying, the product of total steroidal saponins of *Dioscorea nipponica* Makino was obtained with a yield of 0.56% and the content thereof was 95 % (w/w).

[0036] Example 4 Preparation and identification of the compounds in the pharmaceutical composition of the present invention:

1278 g crude extract obtained in example 1 was taken and dispersed in 11100 ml water. Extraction was made in turn with acetic ether (7500ml×4) and n-butanol (7500ml×5), obtaining 102 g extract of acetic ether and 503 g extract of n-butanol. 236 g of the part of n-butanol extract was taken for silica gel column chromatography (10.5×52cm, 2000g) and then gradiently eluted with chloroform-methanol-water (9: 1: 0.1). Each flask recovering 500 ml eluent, total 160 parts were collected. Compound 8 (140mg) was separated out with Fr.16-21, and compound 7 (427mg) was separated out with Fr.41-45, and compound 6 (30g) with Fr. 56-65, compound 5 (8.62 g) with Fr. 94-104, which is the steroidal saponin (III) of the present invention. Fr.113-127 was combined as E (26g), which was separated with silica gel columns (6.5×71cm, 800g) and eluted with chloroform-methanol (5:1) saturated with water. With E passing through the column, Fr.14-20 was combined with as B (4 g), and after ODS purification compound 4 (1.3 g) was obtained. Fr.128-143 was merged as F (63 g), which was separated with silica gel columns (7×52cm, 850 g) and eluted with chloroform-methanol (5:1) saturated with water. With F passing through the column, Fr. 36-60 was combined with as A (10 g) and compound 3 (115 mg) was obtained after ODS purification. With F passing through the column, Fr.61-68 was combined with as B (11 g) and compound 9 (105 mg) was obtained after ODS purification, which was the steroidal saponin (II) of the present invention. With F passing through the column, Fr.69-78 was combined with as C (10 g) and compound 2 (2.89 g) was obtained after ODS purification, which was the steroidal saponin (I) of the present invention.

[0037] The silica gel used for column chromatography (160-200 M, 200-300 M) and GF254 silica gel used for TLC are the products of Qingdao Oceanic Chemical Factory. The reverse phase silica gel ODS (Cosmosil175 C₁₈-OPN) is the product of Japan Nacalai Tesque Company and the reverse phase silica gel plate RP-18F₂₅₄ is the product of Merck.

[0038] With modern method of spectrum of MS, ¹HNMR, ¹³C NMR, DEPT, HMQC and HMBC and in combination with physical and chemical parameters, the structures of the compounds were identified as follows:

[0039] The mass spectrometer used was Finnigan LCQ^{DECA}, and the nuclear magnetic resonance analyzer in use was Bruker AM-400 with TMS as internal standard.

Compound 9 (pseudoprotogracillin)

[0040] ¹³C NMR data of compound 9 was compared with those of pseudoprotodioscin^[3] and gracillin^[4]. The results showed that the sugar signal binding to 26-C of compound 9 and the chemical shift in aglycon part thereof were totally in accord with those of pseudoprotodioscin, and the extra-sugar signals accorded completely with those of gracillin. Accordingly, it was judged primarily that compound 9 was the product of protogracillin dehydrating one H₂O molecule at C₂₀~C₂₂ (namely pseudoprotogracillin (II)) in the spectrum of HMBC, 1-H-glc''' (δ_H4.83) is correlated with aglycon 26-C (δ_C 74.7), 1-H-rha'' (δ_H 6.40) and 1-H-glc'' (δ_H 5.10) with 2-C-glc' (δ_C 76.8) and 3-C-glc' (δ_C 89.3), respectively, and 1-H-glc' (δ_C 4.95) with 3-C (δ_C 77.7). All this also proved the above-presumed binding position and sequence of sugar.

[0041] In summary, the structure of compound 9 is identified as 26-O-β-D-glucopyranosyl-3β, 26 glyco-25(R)-Δ^{5,20(22)}-diene-furo-3-O-[α-L-Rhamnopyranosyl(1→2)]-β-D-glucopyranosyl(1→3)]-β-D-glucopyranoside, namely pseudoprotogracillin (II).

[0042] Compound 8 (trillin): White needle crystals. ESIMS *m/z*: 577[M+H]⁺, 415[M-glc+H]⁺, 397[M-gle-H₂O+H]⁺, 1151[2M-H]⁺, 575[M-H]⁺; ¹HNMR (400 MHz, C₅D₅N): δ_H1.11(3H, d, CH₃-21, *J*=6.8 Hz), 0.87(3H, s, CH₃-19), 0.80 (3H, s, CH₃-18), 0.67 (3H, d, CH₃-27, *J*=4.6 Hz) present 4 signals of methyl proton of the aglycon of dioscin. ¹³C NMR(100 MHz, C₅D₅N) is shown in Table 1. The aglycon part was compared with that of dioscin, wherein in C-3, δ_C71.7→78.6, in C-2, δ_C83.1.4→30.2, in C-4, δ_C42.3→39.3, all illustrate that C-3 part of aglycone is bound to sugar. The ¹³C NMR data of compound 8 (as shown in Table 3) is in accordance with those reported in reference 1 and 2.

[0043] Compound 7 (Progenin II: 3-O-[α-L-rhamnopyranose (1→4)]-β-D-glucopyranose-dioscin): White needle crystals. ESI-MS *m/z*: 723 [M+H]⁺, 577 [M-rha+H]⁺, 415[M-rha-glc+H]⁺, 397 [M-rha-glu-H₂O+H]⁺, 721 [M-H]⁺, 575 [M-rha-

H]; ¹H NMR (400 MHz, C₅D₅N): δ1.11(3H, d, CH₃-21, *J*=6.9 Hz), 0.88 (3H, s, CH₃-19), 0.80 (3H, s, CH₃-18), 0.66 (3H, d, CH₃-27, *J*=5.2 Hz) present 4 signals of methyl proton of the aglycon of dioscin. The anomeric proton δ4.94 (1H, d, *J*=7.7 Hz) of glucose shows that it belongs to β type and the anomeric proton 6.54(1H, br, s) of rhamnose shows that it belongs to α type. The ¹³C NMR data of compound 7 (as shown in Table 3) is in accordance with those reported in reference 1 and 2.

[0044] Compound 6 (Progenin III: 3-O -[α-L-rhamnopyranose (1→2)]-β-D-glucopyranose-dioscin): White powder. ESI-MS *m/z*: 723 [M+H]⁺, 577 [M-rha+H]⁺, 415[M-rha-glc+H]⁺, 397 [M-rha-glc-H₂O+H]⁺; ¹H NMR (400 MHz, C₅D₅N): δ1.11(3H, d, CH₃-21, *J*=7.0 Hz), 1.02 (3H, s, CH₃-19), 0.80 (3H, s, CH₃-18), 0.66 (3H, d, CH₃-27, *J*=5.0 Hz) present 4 signals of methyl proton of the aglycon of dioscin. δ6.36 (1H, br, s) of rhamnose shows that it belongs to α type. δ5.04 (1H, d, *J*=9.0 Hz) of glucose shows that it belongs to β type. The ¹³C NMR data of compound 6 (see Table 3) is in accordance with those reported in reference 1 and 2.

[0045] Compound 5 (dioscin): White needle crystals. ESI-MS *m/z*: 891[M+Na]⁺, 869[M+H]⁺, 577[M-rha-rha+H]⁺, 437 [M-rha-rha-gle+Na]⁺, 903[M+C1]⁺, 867[M-H]⁻, 721[M-rha-H]⁻; ¹H NMR(400 MHz, C₅D₅N): δ1.74 (3H, d, *J*=6.5 Hz), 1.60 (3H, d, *J*=6.2 Hz) present 2 signals of methyl proton of rhamnose, respectively. δ6.37 (1H, br, s), 5.83 (1H, br, s) shows these two rhamnoses are of α type. δ4.92(1H, d, *J*=6.4 Hz) shows the glucose is of β type. δ1.11 (3H, d, CH₃-21, *J*=6.8 Hz), 1.02(3H, s, CH₃-19), 0.79 (3H, s, CH₃-18), 0.66 (3H, d, CH₃-27, *J*=4.9 Hz) present 4 signals of methyl proton of the aglycon of dioscin, respectively. The ¹³C NMR data of compound 5 (see Table 4) is in accordance with those reported in reference 1, and is dioscin.

[0046] Compound 4 (3b, 26-diol-(25R)-Δ^{5,20(22)}-diene-furo-26-0-β-D-glucopyranoside): White powder. ESI-MS *m/z*: 907 [M+Na]⁺, 885 [M+H]⁺, 723 [M-glc+H]⁺, 577 [M-glc-rha+H]⁺, 437 [M-glc-rha-glc+Na]⁺, 883 [M-H]⁻; ¹H NMR(400MHz, C₅D₅N): δ1.61(3H, s, CH₃-21), 0.99(3H, d, CH₃-27, *J*=6.4 Hz), 0.89(3H, s, CH₃-19), 0.70(3H, s, CH₃-18) present 4 signals of methyl proton of the aglycone of dioscin. δ4.95(1H, d, *J*=7.5 Hz), 4.82(1H, d, *J*=6.1 Hz) show that the glucose in compound 4 is of β type and δ 5.90(1H, br, s) show that the rhamnose of the same is of α type. In the ¹³C NMR data of compound 4, aglycon part was compared with that of compound 7, wherein in C-3, δ71.2→78.3, demonstrating, besides C-26 is bound to sugar, C-3 is also bound to sugar. The ¹³C NMR data of compound 4 is in accordance with those reported in reference 4.

[0047] Compound 3(26-0-β-D-glucopyranose-3β, 26-diol-(25R)-Δ^{5,20(22)}-3-0-[(α-L-(25R)-pyranorhamnose(1→4)]-β-D-glucopyranoside)): White powder. ESI-MS *m/z*: 907 [M+Na]⁺, 885 [M+H]⁺, 723 [M-glc+H]⁺, 577 [M-glc-rha+H]⁺, ¹H NMR (600MHz, C₅D₅N): δ1.62 (3H, s, CH₃-21), 1.02 (3H, d, CH₃-27, *J*=6.5 Hz), 0.88 (3H, s, CH₃-19), 0.70(3H, s, CH₃-18) present 4 signals of methyl proton of the aglycone of dioscin. δ1.77(3H, d, *J*=6.0 Hz) shows the signal at the end of methyl proton of rhamnose. The ¹³C NMR data of compound 3 see table 4. The aglycone part is the same as that of compound 8. The NMR data of compound 3 is in accordance with those reported in reference 5.

[0048] Compound 2 (pseudoprotodioscin): White powder. ESI-MS *m/z*: 1053[M+Na]⁺, 723[M-rha-glc+H]⁺, 577[M-rha-glc-rha+H]⁺, 415[M-rha-glc-rha-glc+H]⁺, 1029[M-H]⁻, 883[M-rha-H]⁻; ¹H NMR(400 MHz, C₅D₅N): δ1.61(3H, s, CH₃-21), 1.03(3H, s, CH₃-19), 0.99(3H, d, CH₃-27, *J*=6.8 Hz), 0.70(3H, s, CH₃-18) present 4 signals of methyl proton of the aglycone of dioscin. δ1.74(3H, d, *J*=7.2 Hz), 1.60 (3H, d, *J*=7.7 Hz) present 2 signals of methyl proton of rhamnose, respectively. δ6.37(1H, br, s), 5.83(1H, br, s) shows these two rhamnoses are of α type. δ4.92(1H, d, *J*=6.9 Hz), 4.81 (1H, d, *J*=7.7 Hz) shows these two glucoses are of β type. ¹³C NMR data of compound 2 (see Table 4) is in accordance with those reported in reference 5.

[0049] Compound 1 (protodioscin): For methods for separation and identification, consulted in citation of Bogang Li; Zhengzhi Zhou. Traditional Chinese Drug Research & Clinical Pharmacology 1994, 13(2): 75-76 can be taken for reference.

Table 1 ¹³C NMR Data Of Compound 2-9 (Solvent: C₅D₅N)

C	9	8	7	6	5	4	3	2
1	37.3	37.4(t)	37.4(t)	37.5(0)	37.5(t)	37.5(t)	37.5(t)	37.5(t)
2	29.9	30.2(t)	30.2(t)	30.1(t)	30.1(t)	30.2(t)	30.2(t)	30.1(t)
3	77.7	78.6(d)	78.3(d)	78.2(d)	77.8(d)	78.3(d)	78.3(d)	78.0(d)
4	38.4	39.3(t)	39.3(t)	38.9(t)	38.9(t)	39.3(t)	38.9(t)	38.9(t)
5	140.5	140.9(s)	140.9(s)	140.8(s)	140.8(s)	140.9(s)	140.8(s)	140.8(s)
6	121.6	121.7(d)	121.7(d)	121.7(d)	121.7(d)	121.7(d)	121.7(d)	121.8(d)
7	32.2	32.2(t)	32.2(t)	32.3(t)	32.3(t)	32.3(t)	32.4(t)	32.4(t)
8	31.2	31.6(d)	31.6(d)	31.6(d)	31.6(d)	31.4(d)	31.4(d)	31.4(d)
9	50.1	50.3(d)	50.3(d)	50.3(d)	50.3(d)	50.3(d)	50.2(d)	50.3(d)
10	36.9	37.0(s)	37.0(s)	37.1(s)	37.1(s)	37.0(s)	37.1(s)	37.1(s)
11	21	21.1(t)	21.1(t)	21.0(t)	21.1(t)	21.1(t)	21.2(t)	21.2(t)

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(continued)

	C	9	8	7	6	5	4	3	2
5	12	39.4	39.9(t)	39.8(t)	39.8(t)	39.8(t)	39.6(t)	39.6(s)	39.6(t)
	13	43.2	40.4(s)	40.4(s)	40.4(s)	40.4(s)	43.4(s)	43.4(s)	43.4(s)
	14	54.7(d)	56.6(d)	56.6(d)	56.6(d)	56.6(d)	54.9(d)	54.9(d)	54.9(d)
	15 5	34.3	32.2(t)	32.2(t)	32.2(t)	32.2(t)	34.4(t)	34.5(t)	34.5(t)
	16	84.3	81.1(d)	81.1(d)	81.1(d)	81.1(d)	84.4(0)	84.4(d)	84.4(d)
10	17	64.3	62.9(d)	62.9(d)	62.8(d)	62.9(0)	64.5(d)	64.4(0)	64.5(d)
	18	13.9	16.3(q)	16.3(q)	16.3(q)	16.3(q)	14.1(q)	14.1(q)	14.1(q)
	19	19.2	19.4(q)	19.4(q)	19.3(q)	19.3(q)	19.4(q)	19.4(q)	19.4(q)
	20	103.4	42.0(d)	41.9(d)	41.9(d)	41.9(0)	103.5(s)	103.6(s)	103.5(s)
	21	11.6	15.0(q)	15.0(q)	14.9(q)	15.0(q)	11.8(q)	11.8(q)	11.8(q)
15	22	152.2	109.2(s)	109.3(s)	109.2(s)	109.2(s)	152.4(s)	152.3(s)	152.4(s)
	23	31.2	31.8(t)	31.8(t)	31.8(t)	31.8(t)	31.4(t)	31.4(t)	31.4(t)
	24	23.5	29.2(t)	29.2(t)	29.2(t)	29.2 (t)	23.7(t)	23.6(t)	23.7(t)
	25	33.7	30.6(d)	30.6(0)	30.5(0)	30.5(d)	33.5(d)	33.5(d)	33.4(0)
	26	74.7	66.8(t)	66.8(t)	66.8(t)	66.8(t)	74.9(t)	74.9(t)	74.9(t)
20	27	17.1	17.3(q)	17.3(q)	17.2(q)	17.2(q)	17.3(q)	17.3(q)	17.3(q)
	C3 Sugar Art Glyc 1'	99.7	102.6(d)	102.7(d)	100.3(d)	100.2(d)	102.7(0)	100.3(d)	100.2(d)
	(inner)2'	76.8	75.4(0)	75.5(d)	79.6(d)	78.1(d)	75.5(d)	79.6(d)	78.6(d)
	3'	89.3	78.5(d)	76.7(d)	77.8(d)	76.8(d)	76.7(0)	77.8(d)	76.8(d)
	4'	71.3	71.7(0)	78.2(d)	71.6(d)	78.7(d)	78.5(0)	71.7(d)	78.6(d)
25	5'	78.3	78.1(d)	77.1(d)	77.9(0)	77.9(d)	77.1 (d)	77.7(d)	77.7(d)
	6'	62.2	62.9(t)	61.5(t)	62.6(t)	61.3(t)	61.5(t)	62.6(t)	61.3(t)
	rha 1"	102		102.4(d)		102.0(d)	102.4(d)		102.0(0)
	(1→4)2"	72.5		72.8(d)		72.4(d)	72.8(d)		72.5(d)
	3"	72.2		72.6(d)		72.8(d)	72.6(d)		72.8(0)
30	4"	73.9		74.0(d)		74.1(d)	74.0(d)		74.1(d)
	5"	69.3		70.3(d)		69.4(d)	70.3(d)		69.5(d)
	6"	18.5		18.5(q)		18.6(q)	18.5 (q)		18.6(q)
	rha 1'''	104.3			102.0(d)	102.9(d)		102.0(0)	102.9(d)
	(1→2)2'''	74.7			72.5(0)	72.4(d)		72.5(d)	72.5(d)
35	2'''								
	3'''	78.3			72.8(d)	72.7(d)		72.8(d)	72.7(d)
	4'''	71.3			74.1(d)	73.8(d)		74.1(0)	73.9(d)
	5'''	77.4			69.4(d)	70.4(0)		69.5(d)	70.4(d)
	6'''	62.2			18.6(q)	18.4(q)		18.7(q)	18.4(q)
40	C-26 Sugar Art glc 1'''	104.7					104.9(d)	104.9(d)	104.9(d)
	2'''	75					75.2(d)	75.2(d)	75.2(d)
	3'''	78.5					78.6(d)	78.6(d)	78.5(0)
	4'''	71.5					71.7(0)	71.7(d)	71.7(d)
	5'''	78.4					78.2(d)	78.3(0)	77.9(d)
50	6'''	62.6					62.9(0)	62.8(0)	62.8(0)

[0050] With the above methods, all compounds in the pharmaceutical composition of the present invention can be separated and identified.

55 Example 5 Quality control of the pharmaceutical composition of the present invention

[0051] 9 reference substances that have been confirmed with their structures were introduced in way of mixed sampling

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for HPLC assay under chromatographic condition as follows: chromatographic column: Alltima C18 4.6×250 mm, 5 m, gradient elution; determined with an evaporative light scattering detector; drift tube temperature: 100°C; gas flow rate: 2.0L / min. The chart of gradient elution is as follows:

Table 2 Gradient elution chart

	0(min)	30(min)	50(min)	60(min)	80(min)
Acetonitrile	15 %	40 %	40 %	100 %	100 %
Water	85%	60%	60%	0	100%

[0052] Chromatogram is shown in Figure 1.

[0053] After HPLC assay for the compositions obtained in example 1, 2 and 3, chromatograms are shown in figure1, figure2, figure3. Two points external standard method of logarithmic equation was applied for calculation of the percentage of the weight of compound I, II and III, as shown in Table 3.

Table 3 Percentage of compound I, II, and III by weight in total steroidal saponins.

	Compound (I) %	Compound (II) %	Compound (III) %
Example 1	59.8	4.7	5.7
Example 2	63.8	4.8	6.8
Example 3	58.4	6.8	9.5

[0054] After follow-up determination of 20 batches of the samples of the pharmaceutical composition of the present invention, it was demonstrated that contents for three compounds thereof were stable relatively, and thus they were made as the indexes for quality control and had rather strong controllability. As shown in Table 4:

Table 4 Percentage of each steroidal saponin of the composition of the present invention by weight to the total steroidal saponins

Batch No.	Protodioscin (%)	Pseudoprotodioscin (%)	Pseudoprotograccillin (%)	Compound 3	Dioscin (%)
1	2.1	60.6	6.2	8.1	7.8
2	0.8	60.2	5.3	9.4	9.3
3	1.6	61.8	5.9	8.4	7.0
4	3.9	58.4	6.8	6.8	9.5
5	1.5	66.6	6.8	1.6	6.8
6	9.8	63.0	4.7	0.86	6.2
7	7.7	63.8	4.8	1.0	6.8
8	3.0	65.7	5.6	0.9	8.4
9	8.6	63.2	4.9	1.1	6.3
10	6.3	64.8	5.4	1.1	6.3
11	11.2	61.2	5.2	1.2	5.8
12	9.8	60.6	6.1	1.2	7.2
13	8.7	62.2	5.3	1.1	7.2
14	2.2	67.7	5.6	1.1	6.5
15	7.7	63.5	5.0	1.1	6.8
16	13.4	59.8	4.7	1.4	5.7
17	2.0	68.6	5.8	1.0	5.4
18	3.3	66.9	4.9	1.7	6.4
19	3.2	66.0	4.7	1.6	8.1
20	6.5	62.4	4.5	1.5	7.2
21	10.1	60.2	5.3	1.3	7.1
23	6.7	65.8	4.5	0.9	5.7
24	2.9	68.1	5.0	1.1	5.9

[0055] It was proved in the above detection that the pseudoprotodioscin (I), pseudoprotogracillin (II), dioscin (III) were all in constant proportion in the pharmaceutical composition of the present invention. By controlling directly the contents of these three compounds, the object for controlling the quality of medicine of the present invention is obtainable. In different raw materials and procedures to extract the same, the pseudoprotodioscin (I) and the protodioscin (IV) in the above table could converse to each other. Nevertheless, tests proved that in the same batch of raw material and with the same method for preparation, the contents of pseudoprotodioscin (I) and the protodioscin (IV) were basically constant and so were the contents of pseudoprotogracillin (II) and protogracillin (V). One of the compounds or two of the same with higher contents could be used as the one for quality control. That is to say the pseudoprotodioscin (I) and/or protodioscin (IV), pseudoprotogracillin (II) and/or protogracillin (V), dioscin (III) were all in constant proportion and by directly controlling the contents of these three compounds, the object for controlling the quality of the medicine of the present invention could be achieved.

Example 6 Preparation of the capsules containing the pharmaceutical composition of the present invention

[0056] Ingredients of the capsule (for 1000 capsules):

Total steroidal saponin (obtained in example 2)	100 g
Starch	150 g

[0057] Steroidal saponin and starch were taken according to the proportion as specified above. After mixing, the drug was encapsuled, each containing 250 mg, and prescribed orally for treatment of cerebrocardiovascular diseases at the dosage of 1~2 caps, 3 times daily for two months.

[0058] The steroidal saponin contained in each capsule shall not be less than 65.0 mg, being calculated by the content of dioscin.

Example 7 Preparation of tablets

[0059] Ingredients of tablet (for 1000 tablets):

Total steroidal saponin (obtained as in example 2)	100 g
HPMC _{LV100}	30g
Lactose	70 g
Magnesium stearate	1 g

[0060] The total steroidal saponin, HPMC and lactose were mixed up and made with 75% of ethanol as the adhesive into wet grains, which were screened with 22 M. After being dried at 50°C for 3 hrs, the grains were again screen at 22 M before being added with magnesium stearate and made into tablets, each weighing 0.15 g. The medicine can be prescribed orally for cerebrocardiovascular diseases at the dosage of 1~2 tabs, 3 times daily for two months.

[0061] Each tablet contains steroidal saponin not less than 65.0 mg when calculated by content of dioscin.

Example 8: Preparation of injections

[0062] Ingredients for injection (for 1000 ml):

Total steroidal saponin	50 g
Tween 80	10 ml
Sodium chloride	8 g

[0063] Total steroidal saponin was added with 10% Na₂CO₃ with pH value adjusted to 7.0~7.5. After refrigeration and filtration, tween 80 and NaCl were added and injection water was added to 1000 ml. After being filtered with G₃ sintered filter funnel (Glass), and sub-packed and then filled and sealed. The final product was obtained after wet sterilization at 100°C. The injection can be prescribed hypodermically for treatment of cerebrocardiovascular diseases at the dosage of 1~2 ml, twice daily for two months.

[0064] Each contains steroidal saponin not less than 65.0 mg, as calculated with content of dioscin.

[0065] Soft capsules and dripping pills of the same could be prepared according to routine method with addition of substrate in common usage.

[0066] The advantageous results of the pharmaceutical composition of the present invention were proved in following pharmacodynamic tests.

Test 1 Effect of pseudoprotogrillin (II) on acute myocardial infarction in rats:

[0067] Pseudoprotogrillin (II) obtained in example 4 was applied in relevant pharmacological tests as follows:

[0068] Wistar rats, each weighing 180~220 g, were randomized into 3 groups, namely control group, pseudoprotogrillin group of high dose and pseudoprotogrillin group of low dose, each of which had 12 rats. The rats in the control group were administered by gastric infusion of distilled water. The other rats in test groups were administered by gastric infusion of the relevant medicinal liquid (prepared with distilled water), according to the high dose (3 mg/kg) or low dose (1.5 mg/kg) listed in following table for each group, respectively. Infusion was performed once a day and continued for one week. 1 hour after last administration, urethane of 1 g/kg was given to the rats for abdominal anesthesia. The animal was fixed on its back and its skin was cut open along the midline of its sternum, along the left edge of which an incision was made for the heart to be squeezed out quickly. Ligation was immediately made at the root of the left anterior descending branch of its coronary artery before putting the heart back into its thorax, which was then closed by suture and spontaneous breathing was restored. The test was ended 3 hours after ligation and the heart was sliced transversely below the line of ligation. 5 slices were blue-stained with nitroterazolium and an multiple color pathological analyzer (MPIAS-500) was applied to measure the areas of normal myocardium and infarct myocardium at a fixed imaging distance and to observe the degree of myocardial infarction. After being treated statistically (t test), the results are shown in Table5.

Table 5 Effect of pseudoprotogrillin (II) on myocardial infarction induced by ischemia of myocardium ($\bar{X} \pm s$)

Group	Dosage mg/kg	Area of infarction mm ²	Weight of infarction g	Percentage of infarct area to ventricle (%)	Percentage of infarct area to the heart (%)
Control		112.23±9.46	0.301±0.055	32.9±3.1	128.6±1.5
High dose	3	380.56±7.69**	0.227±0.032**	24.5±2.6**	20.74±1.9**
Low dose	1.5	91.3±11.49*	0.241±0.045*	28.1±1.9*	25.2±2.3*

In comparison with control group: *P<0.05 **P<0.01

[0069] The results in Table 4 demonstrated that in the groups treated with compound (II) at both high dose or low dose, the area and weight of infarct myocardium, the percentages of infarct area to ventricle and to the whole heart are significantly lower in comparison with the control group (P<0. 01 and P<0. 05, respectively), and present a dose-effect relationship, indicating the novel compound pseudoprotogrillin (II) of the present invention can minimize the extent of myocardial infarction, reduce the area of infarction and weight of infarct area and has obvious action to protect myocardium.

[0070] The above pharmacodynamical test does not only demonstrate that the compound pseudoprotogrillin (II) in the pharmaceutical composition of the present invent is a novel compound, but also proves strongly that the new compound can be prepared independently into an agent with definite pharmacodynamic action.

Test 2 Effects of the pharmaceutical composition of the present invention on angina, blood lipids and platelet aggregation

[0071] 1084 patients with thoracic obstruction (angina) were clinically observed in a way of random, double blind and control. They were prescribed at random with A1, A2, A3, A4, Fufangdanshen Tablets and isosorbide dinitrate.

[0072] All cases to be administered with the drugs had gone through tests of proportionality (comparability) in aspects of genders, ages, courses of disease and complications. The results proved that the patients in groups taking A1, A2, A3 and A4 capsules and in group taking Fufangdanshen Tablet and isosorbide dinitrate had comparability with no obvious differences in their genders, ages, courses of disease and complications.

[0073] 6 drugs labeled as A1, A2, A3, A4, B and C were all encapsuled with same appearance. Five of the drugs were taken orally, 1 capsule, 3 times daily. These 5 drugs had 8 weeks as one therapeutic course. During the period for administration of these 6 drugs, all patients stopped taking any medicine that might have effects on their coronary blood flow, blood lipids and blood pressure. In case of emergency, the said medicine could be applied temporarily, but once the condition turned better, the said medicine should be withdrawn and the dose should be recorded.

[0074] Unblinding: A1, A2 and A3 were the capsules comprising the pharmaceutical composition obtained in example 1, example 2 and example 3 and prepared with the method in example 6. A4 was the capsules comprising the composition extracted with old procedures and prepared with the method in example 6. The content of total steroidal saponin in each capsule of A1, A2, A3 and A4 was the same. The dose was 200 mg each time, 3 times daily. B was Fufangdanshen

Tablet, the dose of which was 3 tabs, 3 times daily. C was the drug of isosorbide dinitrate and the dose for it was 5 mg, 3 times daily.

[0075] All patients should be withdrawn from the medicines that have effects on coronary heart disease 3 days before receiving testing drugs, and should have their blood and urine tests, hepatic and renal function exams, and blood lipids, rest ECG and load test analyzed. After treatment, all of the above said indexes should be rechecked.

[0076] During treatment, all medicines to reduce blood pressure and blood fat and to expand coronary artery should be withdrawn. Only those patients with frequent angina and serious cardiac arrhythmias, nitroglycerin drugs could be prescribed temporarily, and dosage and time of stoppage should be recorded.

[0077] Criteria to assess therapeutic effects: *Standard for Therapy of Coronary Heart Disease and Angina and Electrocardiogram* amended in 1979 in the National Research Symposium on Treatment and Prevention of Coronary Heart Disease, Angina and Arrhythmia with Combination of Traditional Chinese Medicine and Western Medicine was adopted for therapeutic assessment.

[0078] Results are shown in table 6 and 7:

Table 6 Comparison of the effects of 5 drugs on angina

group	Total cases	Obvious effective	%	Effective	%	No effective	%	Total effective%
A1 group	178	72	40.4	86	48.3	20	10.1	89.9
A2 group	179	74	41.3	88	49.2	17	9.5	90.5
A3 group	178	71	39.9	90	50.6	17	9.6	90.4
A4 group	258	99	38.4	117	45.3	42	16.3	83.7
Fufangdanshen	123	34	27.6	44	35.8	45	36.6	63.4
Tablets								
isosorbide dinitrate	168	37	22.0	80	47.6	51	30.4	69.6

[0079] By Ridit analysis, all p values in A1, A2 and A3 groups were less than 0.01 in comparison with those in B group and in C group, respectively. By X^2 analysis, A1, A2 and A3 group presented an obvious effective rate in comparison with B group, $P < 0.05$; A1, A2 and A3 group presented an obvious effective rate in comparison with C group, $P < 0.01$. The results demonstrated that the therapeutic effects of the capsules made from the composition obtained in example 1, example 2 and example 3 of the present invention are better than those of Fufangdanshen Tablet (B) and isosorbide dinitrate (C). The total effective rates to relieve angina were 89.9%, 90.5 % and 90.4 % respectively and definitely, proving that an effective and stable pharmacodynamic actions were achieved by controlling of 3 components.

[0080] In comparison with routine method, the preparation procedures of the present invention decrease cost by more than 35% and increase yield by 13%. With same amount of raw material, the pharmacodynamic action increased by 5% when the pharmaceutical composition of the present invention was adopted in clinical observation of 258 cases.

Effect on ischemia of myocardium

[0081] In 8 weeks for a course of treatment, the comparison of the pharmaceutical composition of the present invention is shown in Table 7.

Table 7 Comparison of therapeutic effects of 6 drugs on myocardial ischemia

group	Total cases	Obvious effective	%	Effective	%	No effective	%	Total effective%
A1 group	132	41	31.1	31	23.5	60	45.4	55.5
A2 group	132	42	31.8	32	24.2	58	43.9	56.0
A3group	131	41	31.1	31	23.7	59	45.0	54.8
A4group	258	76	29.4	58	22.5	124	48.1	51.9
Fufangdanshen	57	10	17.5	12	21.1	35	61.4	38.6
tablets	85	25	29.4	20	23.5	38	44.7	52.9
isosorbide dinitrate								

[0082] By Ridit analysis, the therapeutic effects on ischemia of myocardium in A1, A2 and A3 groups had no obvious differences with that in isosorbide dinitrate group, respectively (all $P>0.05$), but surpassed that in Fufangdanshen Tablet group (all $P<0.01$). By X^2 analysis, the obvious effective rates of A1, A2 and A3 groups were higher than that in Fufangdanshen group, $P<0.05$, but with no differences with that in isosorbide dinitrate group, all $p<0.05$, indicating that capsules made from the pharmaceutical composition of the present invention obtained as in example 1, example 2 and example 3 had better therapeutic effects on angina than that of Fufangdanshen Tablet (B), and had the equivalent effect as that in isosorbide dinitrate group (C). The total effects on ST-T segments of the rest ECG were 55.5%, 56.0 % and 54.8% respectively, proving the effects were definite and effective and stable pharmacodynamic actions had been achieved by controlling 3 components. The effects of the pharmaceutical composition of the present invention were 5% higher than that prepared with old procedures.

Effect on blood lipids

[0083] Among 164 patients with hypercholesteremia and 162 patients with hypertriglyceridemia before treatment, 78 patients with hypercholesteremia turned normal and so did 87 patients with hypertriglyceridemia after treatment. The cholesterol in their blood dropped from 6.8 ± 1.1 mmol/L before treatment to 6.3 ± 1.1 mmol/L after treatment. The triglyceride in their blood dropped from 2.4 ± 0.8 mmol/L before treatment to 1.9 ± 0.7 mmol/L after treatment. The differences were of great significance ($P<0.01$) by t test.

Effect on platelet aggregation

[0084] The pharmaceutical composition had an obvious effect on platelet aggregation induced by adenosine diphosphate (ADP) and adrenergic. ADP decreased the rate of platelet aggregation from $68\%\pm 15\%$ before being treated with the composition to $60\%\pm 15\%$ after being treated composition; adrenergic dropped the aggregation rate from $73\%\pm 15\%$ before being treated with the composition to $64\%\pm 12\%$ after being treated, indicating a difference of great significance after t test ($P<0.01$).

[0085] Improvement of the pharmaceutical composition of the present invention on clinical symptoms of patients with coronary heart disease

Table 8 Improvement of main symptoms before and after treatment with the capsules comprising the pharmaceutical composition of the present invention

	Total cases	Obviously effective	effective	No effect	Worsen	Rate of obvious effect(%)	Total effective rate (%)
chest distress	244	112	212	12	0	46	87
cardiopalmus	225	123	193	31	1	55	86
short breath	263	104	203	59	1	40	77
acratia	226	68	155	69	2	30	68
dizziness	206	83	147	58	1	40	71
headache	168	58	117	49	2	34	70

Adverse effects

[0086] Among 825 cases in this test, 5 had dizziness during the period of administration, 9 had stomach or abdominal discomfort, 2 had headache and 1 had hectic. All of the patients with above-mentioned adverse effect were not affected in their treatment and their routine blood and urine tests, hepatic and renal function exams presented normal before and after treatment.

[0087] It is well illustrated by the above-mentioned preparation procedures, structural determinations, quantified control and pharmacodynamic tests that simultaneous control of the three compounds of the pharmaceutical composition of the present invention, namely pseudoprodigiosin (I), pseudoprotogracillin (II) and diocin (III), and effective quality control thereof can increase the pharmacodynamic action and stability of the drugs made thereof with minimized dosage and convenient usage thereof.

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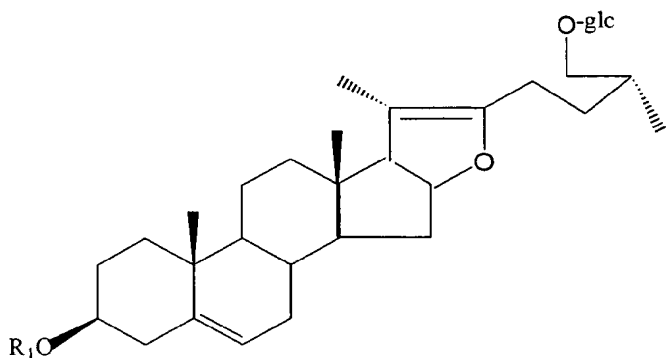
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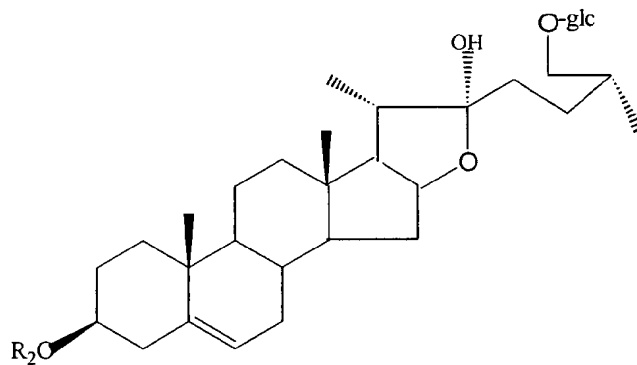
Claims

1. A pharmaceutical composition containing steroidal saponins, comprising:

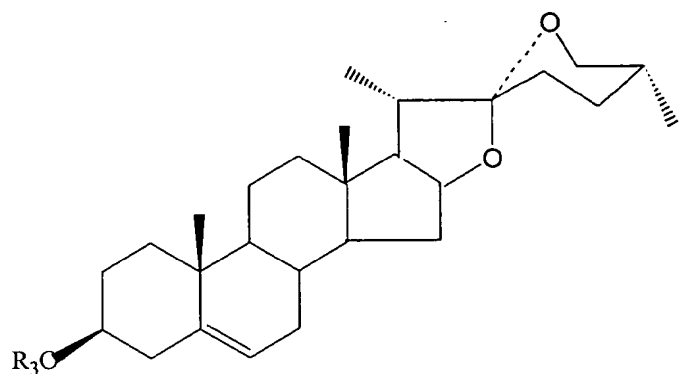
5-25 parts by weight of furostanol saponin represented by general formula A and/or general formula B; and
1-10 parts by weight of spirostanol saponin represented by general formula C,



General Formula A

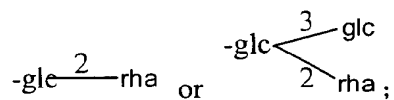
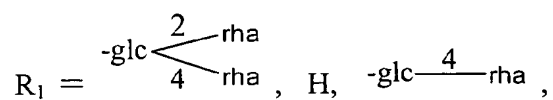


General Formula B

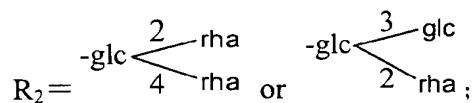


General Formula C

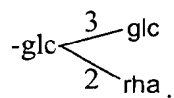
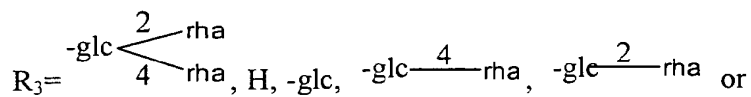
wherein, in general formula A,



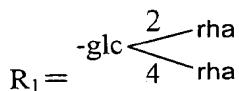
in general formula B:



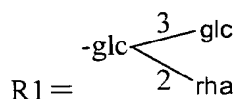
in general formula C:



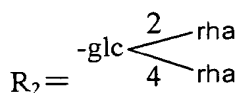
2. The pharmaceutical composition according to claim 1, wherein in general formula A: when



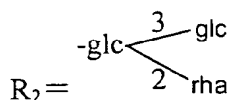
the compound represented by formula A is pseudoprotodioscin (I);
when



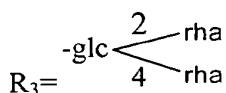
the compound represented by formula A is pseudoprotogracillin (II);
in general formula B, when



the compound represented by formula B is protodioscin (IV);
when



the compound represented by formula B is protogracillin (V);
in general formula C, when



the compound represented by formula C is dioscin(III), and said composition comprises:

5~22 parts by weight of pseudoprotodioscin (I) and/or protodioscin (IV);
1~5 parts by weight of pseudoprotogracillin (II) and/or protogracillin (V); and
1~8 parts by weight of dioscin (III).

3. The pharmaceutical composition according to claim 2, wherein it comprises:

5~20 parts by weight of pseudoprotodioscin (I),
1~3 parts by weight of pseudoprotogracillin (II), and
1~5 parts by weight of dioscin (III).

4. The pharmaceutical composition according to claim 3, wherein it comprises:

12~18 parts by weight of pseudoprotodioscin (I),
1 part by weight of pseudoprotogracillin (II), and
1.2~2.5 parts by weight dioscin (III).

5. The pharmaceutical composition according to any one of claim 1~4, wherein said steroidal saponins are originated from the extracts of *Dioscorea panthaica* Prain et Burkill and *Dioscorea nipponica* Makino, both belonging to species of Dioscorea, Dioscoreaceae.

6. The pharmaceutical composition according to claim 5, wherein in said extract, the content of total steroidal saponin is no less than 65% (w/w) when calculated by dioscin.

7. The pharmaceutical composition according to claim 5 or 6, wherein in said extract, the total contents of three kinds of steroidal saponins, pseudoprodioscin (I), pseudoprotogracillin (II) and discin (III), is no less than 50% (w/w) of the content of total steroidal saponins.

8. The pharmaceutical composition according to claim 5, 6 or 7, wherein said extract has the HPLC fingerprint as shown in Fig 1, in which the characteristic peaks of said HPLC fingerprint are as follows:

the retention time for pseudoprotodioscin (I): 28.27 min;
the retention time for pseudoprotogracillin (II): 29.5 min; and
the retention time for dioscin (I): 57.10 min;
the chromatographic conditions of HPLC are as follows :

chromatographic column: Alltima C18 4.6×250 mm, 5 m;
gradient elution;
determined with an evaporative light scattering detector;
drift tube temperature: 100°C ; and
gas flow rate: 2.0L / min.

9. The pharmaceutical composition according to claim 5~8, wherein said extract is prepared by the method comprising the steps of:

a. taking rootstocks of *Dioscorea panthaica* Prain et Burkill and *Dioscorea nipponica* Makino, or the rootstocks of freshly-collected *Dioscorea panthaica* Prain et Burkill and *Dioscorea nipponica* Makino as raw material;

crushing and slicing the rootstocks;
extracting with one or more than one kinds of solvents selected from a group consisting of water, methanol, ethanol, n-butanol and other low-fat alcohols, the amount of said solvent shall be 15~25 times of the raw material;

b. cooling and filtering the extract prepared in step a;

passing the filtrate through an absorbent resin column,
abandoning the effluent and washing with water till the effluent turns colorless;
abandoning the rinsing water;

c. eluting the absorbent resin column that is washed with water in step b with one or more than one kinds of solvent selected from a group consisting of ethanol, methanol, acetone, 50~90% ethanol, 30~80% hydrous methanol and 60~95% hydrous acetone;

collecting and condensing the effluent;

d. adding 60~95% alcohol to the concentrated solution obtained in step c for alcohol precipitation, filtering and collecting the filtrate;
condensing and drying the filtrate to obtain product.

10. The pharmaceutical composition according to claim 9, wherein in step b, it further includes a step of decompressing condensation before cooling and filtering when the extract contains methanol, ethanol, n-butanol or other low-fat alcohols.

11. The pharmaceutical composition according to any of claim 1~10, wherein it comprises said steroidal saponins or the extracts of *Dioscorea panthaica* Prain et Burkill and *Dioscorea nipponica* Makino as an active ingredient and pharmaceutically acceptable adjuvants.

12. The pharmaceutical composition according to claim 11, wherein said pharmaceutical composition is in form of tablet,

capsule, soft capsule, grain, oral liquor, dripping pill or injection.

13. A method for preparing the pharmaceutical composition according to any of claim 1~12, wherein it includes the following steps:

taking by weighing furostanol saponin of general formula A and/or general formula B and spirostanol saponin of general formula C, mixing them with the ratio by weight of 5~25 parts of furostanol saponin and 1~5 parts of spirostanol saponin; adding pharmaceutically acceptable adjuvants to obtain the pharmaceutical composition.

14. A method for preparing the pharmaceutical composition according to any of claim 1~12, wherein it includes the following steps:

a. taking rootstocks of herbs *Dioscorea panthaica* Prain et Burkill and *Dioscorea nipponica* Makino, or the rootstocks of freshly-collected *Dioscorea panthaica* Prain et Burkill and *Dioscorea nipponica* Makino as the raw material, slicing rootstocks or smashing them, extracting with one or more than one kinds of solvent selected from a group consisting of water, methanol, ethanol, n-butanol and other low-fat alcohols, the amount of which shall be 15~25 times of the raw material;

b. cooling and filtering the extract prepared in step a, passing the filtrate through absorbent resin column, abandoning the effluent and washing with water till the effluent turns colorless; abandoning the rinsing water;

c. eluting the absorbent resin column that was washed with water in step b with one or more than one kinds of solvent selected from a group consisting of ethanol, methanol, acetone, 50%~90% ethanol, 30%~80% hydrous methanol and 60%~95 % hydrous acetone, collecting and condensing the effluent;

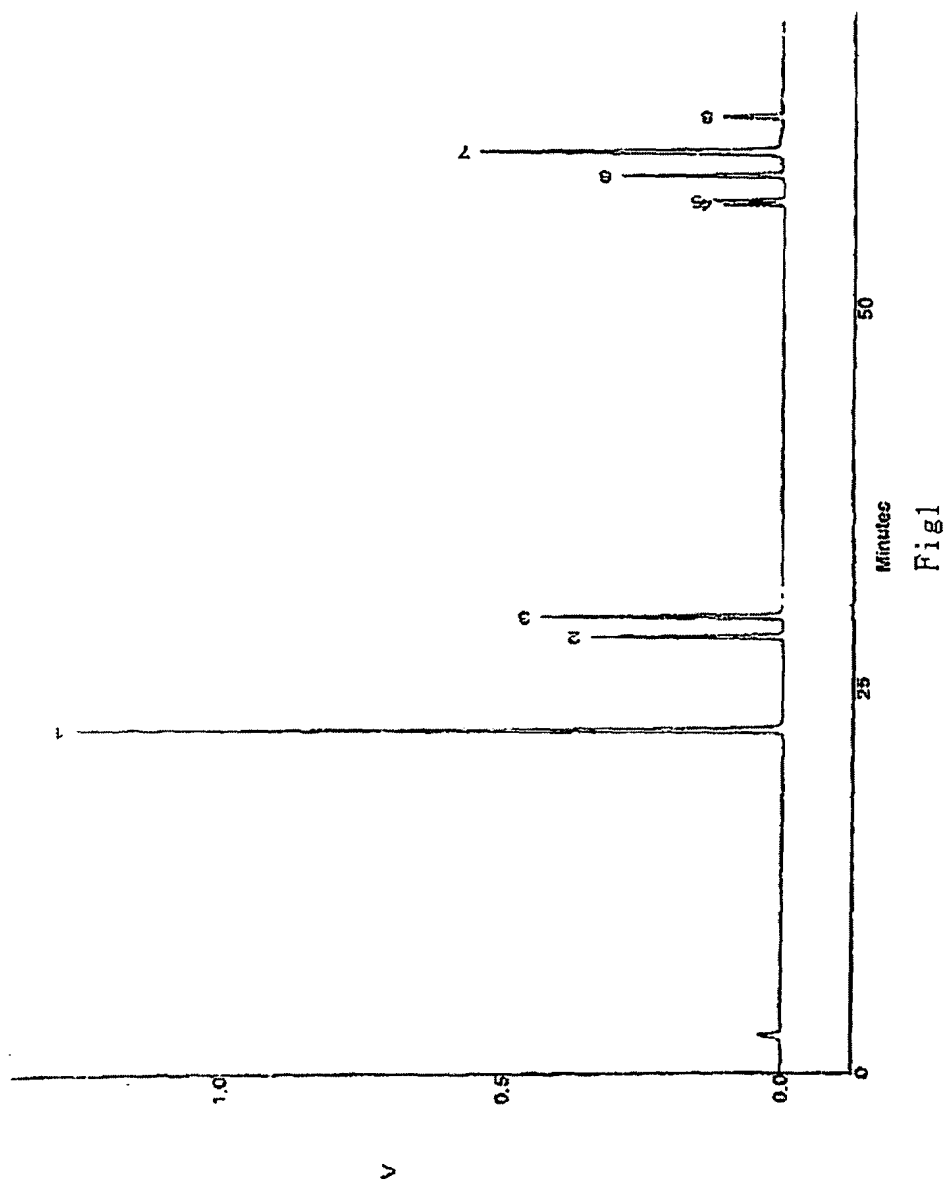
d. adding 60%~95 % alcohol to the concentrated solution obtained in step c for alcohol precipitation, collecting filtrate after filtered;

e. condensing and drying the filtrate in step d, adding pharmaceutically acceptable adjuvants to obtain the pharmaceutical composition.

15. The method according to claim 14, wherein in step b, it further includes a step of decompressing condensation before cooling and filtering when the extract contains methanol, ethanol, n-butanol or other low-fat alcohols.

16. A use of the pharmaceutical composition of claim 1 to 12 in preparing the pharmaceutical for treatment and prevention of cerebrocardiovascular diseases.

17. The use according to claim 16, wherein said diseases are coronary heart disease, angina, myocardial infarction, arrhythmia, hyperlipemia or ischemic cerebrovascular diseases.



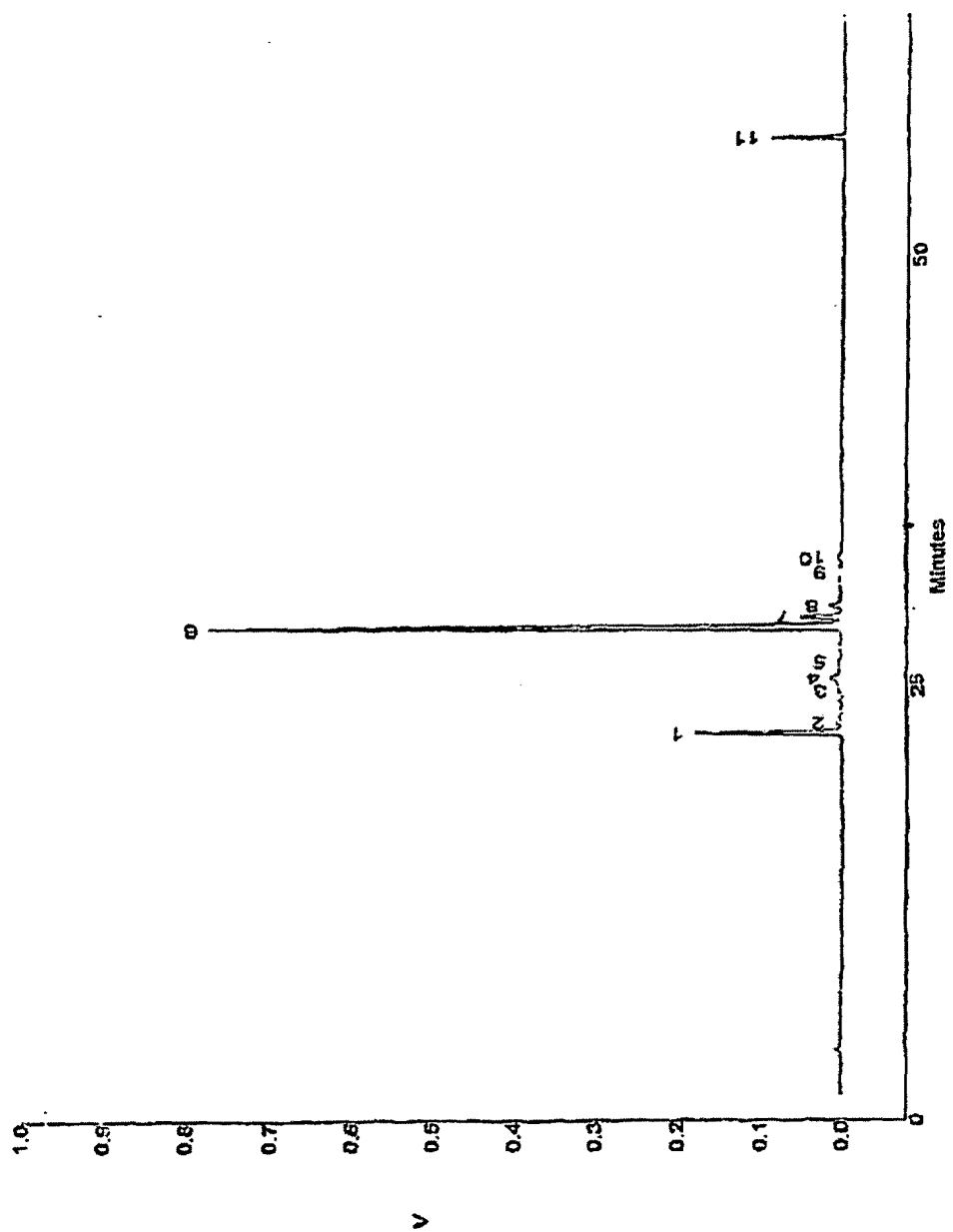


Fig2

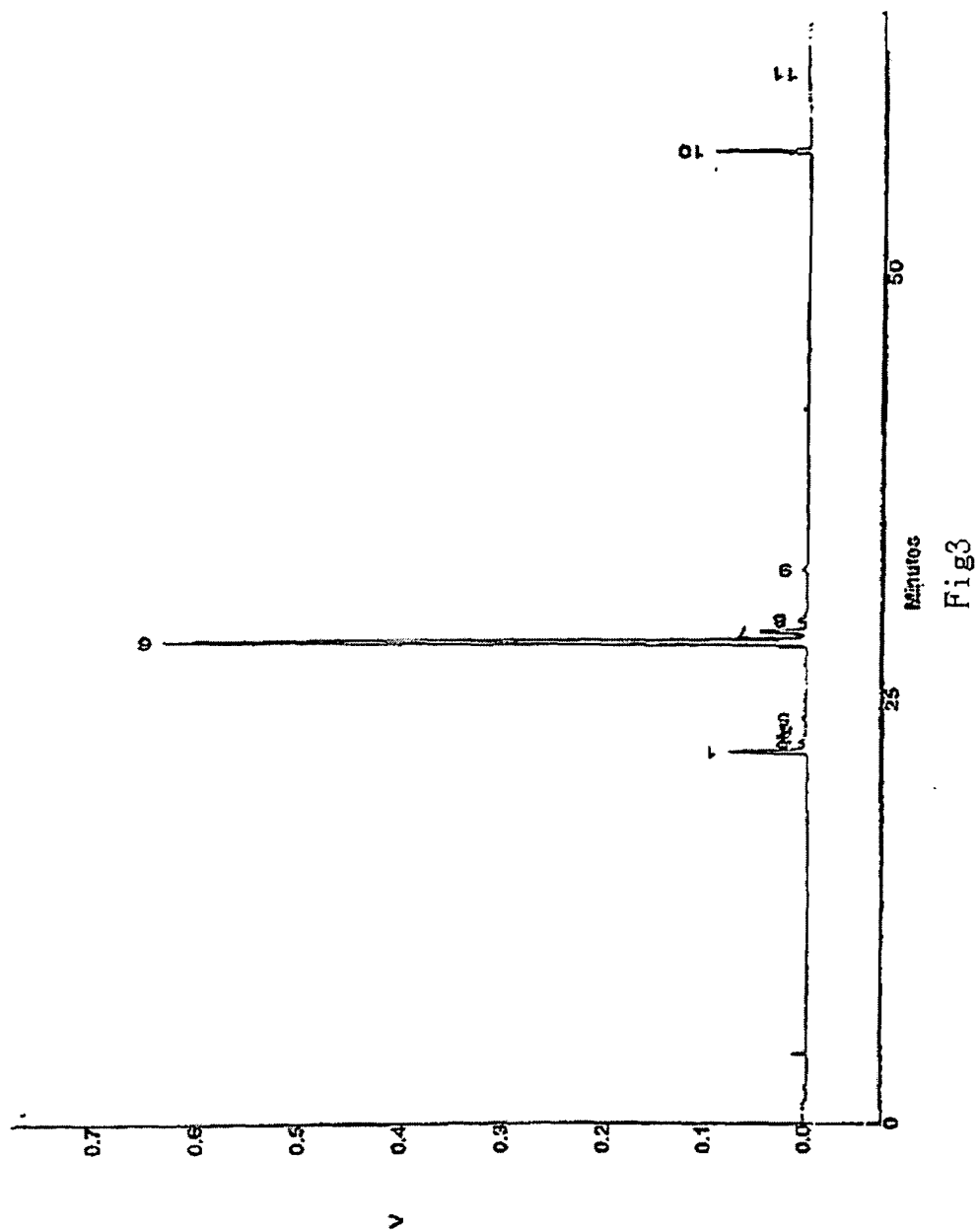
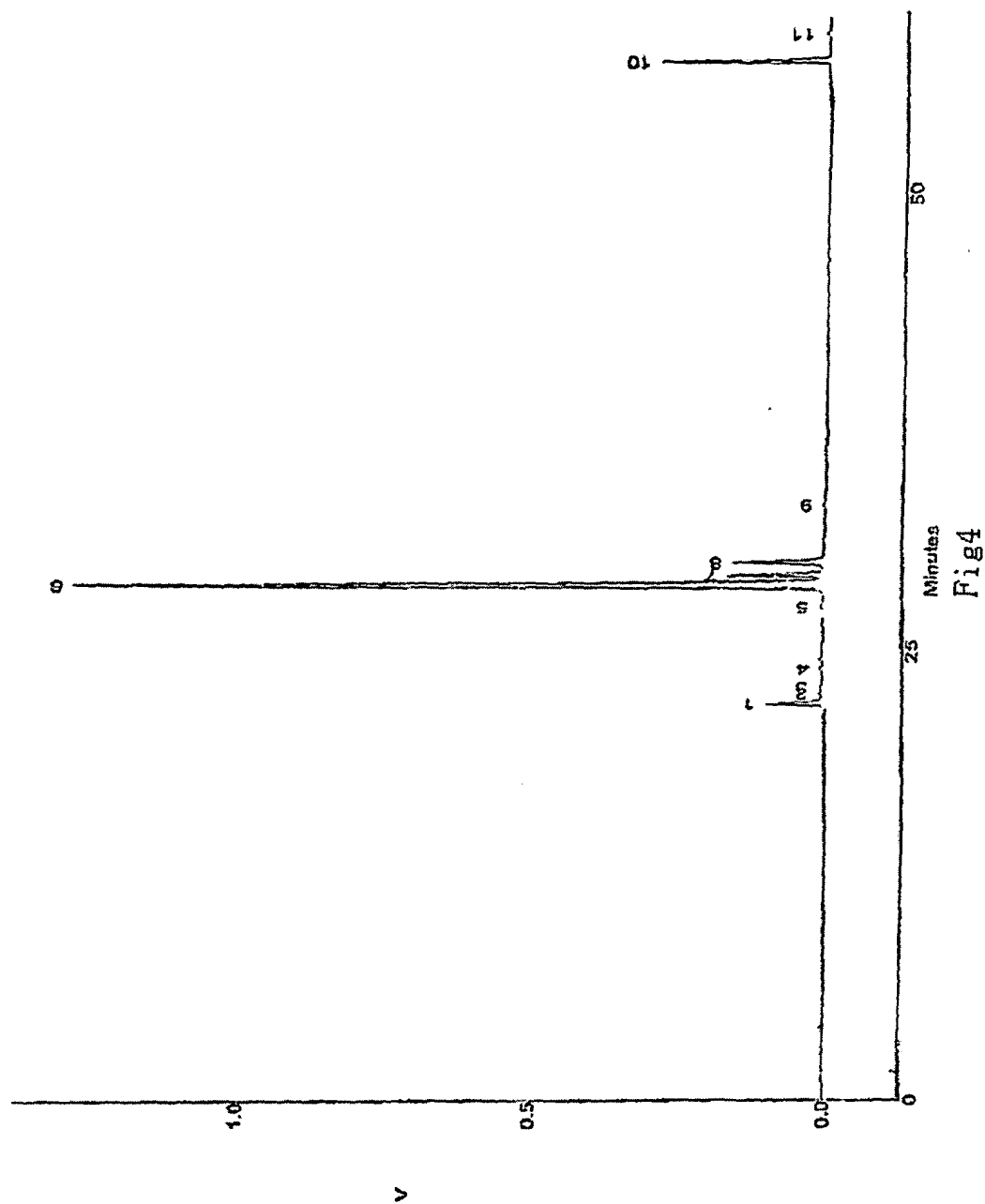


Fig3



INTERNATIONAL SEARCH REPORT

International application No.

PCT/CN2005/001621

A. CLASSIFICATION OF SUBJECT MATTER				
IPC ⁷ A61K 31/7048, A61P 9/10, A61P 9/06, A61P 3/06				
According to International Patent Classification (IPC) or to both national classification and IPC				
B. FIELDS SEARCHED				
Minimum documentation searched (classification system followed by classification symbols)				
IPC ⁷ A61K 31, A61P 9, A61P 3				
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched				
CPRS, CNKI (CN)				
Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)				
WPI, EPODOC, PAJ, BA, MEDLINE, EMBASE, CA dioscin, trillin, gracillin, protodioscin, protogracillin, pseudoprotodioscin, dioscorea panthaica, dioscorea nipponica etc.				
C. DOCUMENTS CONSIDERED TO BE RELEVANT				
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.		
X	NEW DRUGS AND CLINICAL REMEDIES, 1994, 13(2), Li bogang et al. : "The chemistry of	1-7, 11-13		
Y	new drug Di'ao Xinxuekang for treating heart-blood disease." Pp75-76	16-17		
Y	Chinese Pharmacopoeia 2000 th Part 1. Chemical Industry Publishing Company. Pp441-442	16-17		
X	CN1511535 A (CHEN-N) 14.Jul.2004 (14.07.2004) See entire document, especially page 5, line 29 – page 6 line 3.	1-15		
X	CN1389469 A (KUNM-N) 08.Jan.2003 (08.01.2003) Example 1	1-15		
A	CN1415625 A (TIAN-N) 07.May.2003 (07.05.2003) See entire document.	1-17		
A	KR2004077648 A (UYKY-N) 06.Sep.2004 (06.09.2004) WPI / DERWENT abstract	1-17		
<input type="checkbox"/> Further documents are listed in the continuation of Box C. <input checked="" type="checkbox"/> See patent family annex.				
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"A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier application or patent but published on or after the international filing date "L" document which may throw doubts on priority claim (S) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family			
Date of the actual completion of the international search		Date of mailing of the international search report		
23. Dec. 2005 (23.12.2005)		05 · JAN 2006 (05 · 01 · 2006)		
Name and mailing address of the ISA/CN The State Intellectual Property Office, the P.R.China 6 Xitucheng Rd., Jimen Bridge, Haidian District, Beijing, China 100088 Facsimile No. 86-10-62019451		Authorized officer Xiao Ying Telephone No. (86-10) 62085253		

Form PCT/ISA/210 (second sheet) (April 2005)

INTERNATIONAL SEARCH REPORT
Information on patent family membersInternational application No.
PCT/CN2005/001621

Patent Documents referred in the Report	Publication Date	Patent Family	Publication Date
CN1511535 A	2004-07-14	NONE	
CN1389469 A	2003-01-08	NONE	
CN1415625 A	2003-05-07	NONE	
KR2004077648 A	2004-09-06	NONE	

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REFERENCES CITED IN THE DESCRIPTION

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Patent documents cited in the description

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- CN 02112159 [0004]

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